

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 116 774 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
18.07.2001 Bulletin 2001/29

(51) Int Cl.7: **C09K 21/14**

(21) Application number: **00118672.5**

(22) Date of filing: **29.08.2000**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **17.01.2000 JP 2000007515**
09.03.2000 JP 2000064370

(71) Applicant: **SUMITOMO BAKELITE COMPANY
LIMITED**
Tokyo 140 (JP)

(72) Inventor: **Tobisawa, Akihiko**
Fujieda-shi (JP)

(74) Representative: **Vossius, Volker, Dr. et al**
Patent- und Rechtsanwaltskanzlei
Holbeinstrasse 5
81679 München (DE)

(54) **Flame-retardant resin composition, and preregs and laminates using such composition**

(57) The present invention provides a resin composition having excellent flame retardancy without using any halogen-containing flame retardant, and preregs and laminates using such a resin composition. More specifically, the present invention provides a flame-retardant resin composition comprising as essential com-

ponents (A) an epoxy resin, (B) a curing agent and (C) a phosphorus compound containing at least as part thereof a phosphine oxide compound, a prepreg produced by impregnating this flame-retardant resin composition in a fiber base, and a laminate produced by hot-pressing a single sheet or a pile of two or more sheets of the prepreg.

EP 1 116 774 A2

Description

[0001] The present invention relates to a resin composition having excellent flame retardancy with no halogen-containing flame retardant incorporated, and prepregs and laminates using such a resin composition.

[0002] Owing to their surpassing properties, thermosetting resins represented by epoxy resins are prevalently used for electrical and electronic machine parts, and in many cases these resins are provided with flame retardancy for securing safety against fires in use. Conventionally, the resins have been made flame-retardant by incorporating a halogen-containing compound such as brominated epoxy resin. Although such halogen-containing compounds have high flame retardancy, aromatic bromine compounds have the problem that they not only release corrosive bromine and hydrogen bromide in the course of pyrolysis but also have a possibility of forming highly toxicant polybromodibenzofuran and polydibromobenzoxine when decomposed in the presence of oxygen. Further, great difficulty comes in disposing of superannuated wastes containing bromine. For these reasons, study is now being made in earnest on use of phosphorus and nitrogen compounds as well as inorganic fillers as a flame retardant in place of the conventional bromine-containing ones.

[0003] As mentioned above, flame retarding of resins can be realized by making use of a phosphorus compound, a nitrogen compound and an inorganic filler. This is based on the concept that the nitrogen and phosphorus compounds accelerate carbonization of a resin to prevent its combustion, while the inorganic fillers help to reduce the amount of a combustible resin by their addition, consequently making the resin less combustible. Inorganic fillers such as metal hydroxides release water when burned to lower temperature and thereby prevent combustion. It has been known in the art, as disclosed in for instance JP-A-10-195178 and JP-A-10-166501, to render resins flame-retardant by incorporating a phosphorus compound and a nitrogen compound along with an inorganic filler instead of using a halogen compound, and prepregs and laminates using such resin compositions have been put to practical use.

[0004] The circuit-board manufacturing process involves various chemical liquid treating steps. In the desmearing step, for instance, a basic compound such as sodium hydroxide is required, while use of an oxidizing compound is essential for the blackening treatment. Phosphorus compounds such as phosphoric esters, which are commonly used as a flame retardant, have the problem that they tend to react with the above-mentioned treating compounds to give rise to difficulties.

[0005] As a phosphorus compound reactive with epoxy resins and epoxy resin curing agents, other than phosphoric esters, there is known 9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (produced and sold under the trade name of HCA by Sanko Chemical Co., Ltd.). (See, for example, JP-A-11-124489 and JP-A-2000-7898). HCA is hardly hydrolyzable and has excellent chemical resistance, but it is highly reactive with epoxy resins, so that when HCA alone is used as a phosphorus compound, its content must be increased and the reaction proceeds in the state of pre-curing varnish with an epoxy resin, and hence the final product such as a laminate may not be provided with the desired properties such as soldering heat resistance.

[0006] The present invention has been achieved in the course of studies for solving the above problems, and it provides a resin composition having excellent chemical resistance and flame retardancy by using a phosphine oxide compound with a low tendency to hydrolyze as a phosphorus-containing flame retardant. It is also envisaged in the present invention to provide prepregs using the resin composition, and flame-retardant laminates comprising such prepregs.

[0007] Specifically, the present invention provides a flame-retardant resin composition comprising as essential components (A) an epoxy resin, (B) a curing agent, and (C) a phosphorus compound containing at least as a part thereof a phosphine oxide compound. The present invention also provides a prepreg obtained by impregnating the flame-retardant resin composition in a base, and a flame-retardant laminate or a copper-clad flame-retardant laminate produced by hot-pressing a single sheet or a pile of two or more sheets of the prepreg.

[0008] As mentioned above, the added type phosphoric esters generally used as a phosphoric flame retardant, such as triphenyl phosphate and cresyl diphenyl phosphate, are easily hydrolyzed and tend to be eluted in a chemical solution. Therefore, resin compositions comprising such compounds as a flame retardant involve problems in their use for laminates.

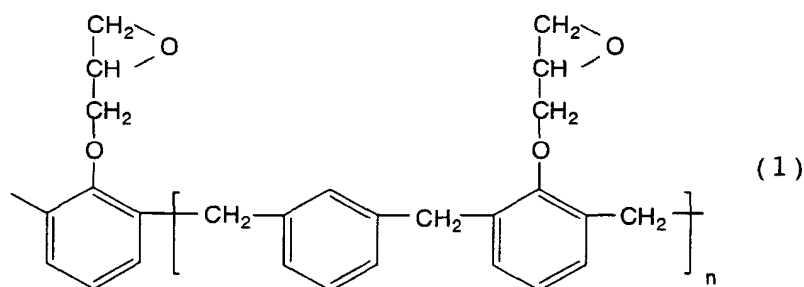
[0009] The phosphine oxide compounds are less susceptible to hydrolysis as they have P-C bonds. These compounds, therefore, unlike the phosphoric esters having P-O bonds, have surpassing chemical resistance. Also, they show satisfactory flame retardancy by addition of only a small quantity because of high phosphorus content.

[0010] Generally, phosphorus compounds are hygroscopic, so that too much loading of a phosphorus compound gives adverse effects to moisture resistance and other properties of the cured resin composition. In contrast, phosphine oxide compounds have a restrained tendency to hydrolyze and low hygroscopicity. It is also notable that use of a novolak epoxy resin and a novolak resin as a curing agent makes it possible to provide satisfactory flame retardancy of the composition by addition of a smaller amount of phosphorous compound than required in the conventional art, because these resins have a high content of benzene rings and are therefore high in heat resistance and flame retardancy.

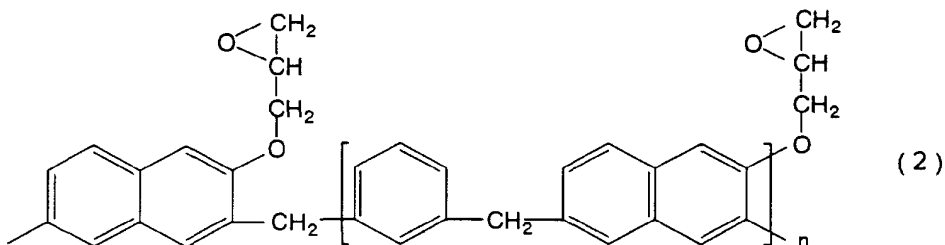
[0011] Thus, the flame-retardant resin composition according to the present invention is characterized by use of an epoxy resin and a curing agent with high heat resistance and use of a phosphine oxide compound as a flame retardant to realize satisfactory flame retardancy and chemical resistance of the composition without using a halogen compound.

[0012] Examples of the epoxy resins (A) usable in the present invention include bisphenol A epoxy resins, bisphenol F epoxy resins, phenolic novolak epoxy resins, cresol novolak epoxy resins, and bisphenol A novolak epoxy resins, and are not limited to these epoxy resins. It is also possible to use two or more different types of epoxy resins simultaneously.

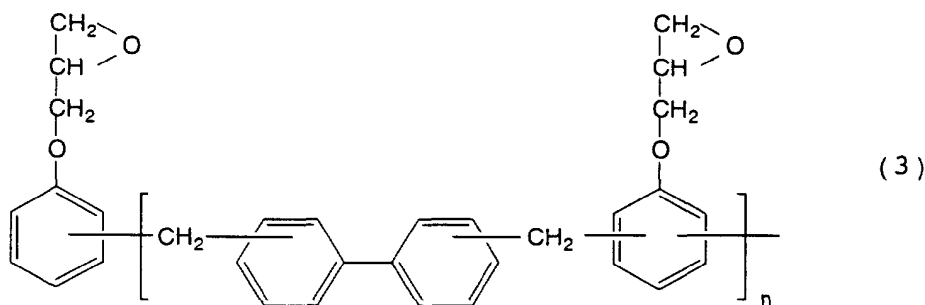
[0013] In view of heat resistance, it is preferred to use the novolak epoxy resins with a high benzene ring content and high heat resistance, especially phenolic novolak epoxy resins and cresol novolak epoxy resins. The phenolic aralkyl epoxy resins represented by the following formula (1), the naphthalene aralkyl epoxy resins represented by the following formula (2) and the biphenyl-modified epoxy resins represented by the following formula (3) are also preferred, as these resins have a high content of aromatic rings and are small in ratio of polar groups, and hence high in heat resistance and capable of reducing water absorption of the composition:



wherein n is a natural number,



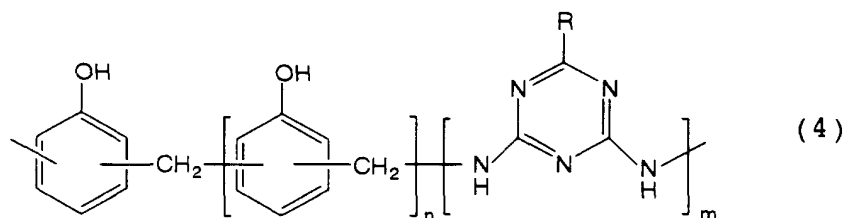
wherein n is a natural number, and



wherein n is a natural number.

[0014] In the present invention, the above type of epoxy resin is preferably used in an amount of at least 50% by weight of the whole epoxy resins for providing the desired improvement of heat resistance and flame retardancy.

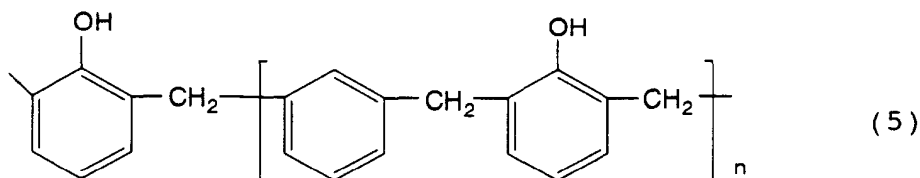
[0015] As the curing agent used as component (B) in the present invention, novolak resins, amine compounds, acid anhydrides and the like can be used. Novolak resins are preferred, as they have high heat resistance and allow reduction of the amount of the phosphorus compound to be added. Phenolic novolak resins and phenolic aralkyl resins are usually used. Use of a triazine-modified novolak resin represented by the following formula (4) is preferred, as flame retardancy is further improved by the presence of triazine ring which is a nitrogenous component:



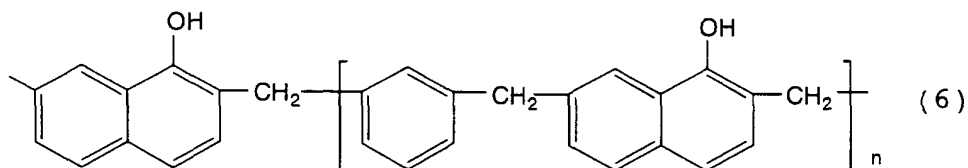
wherein R is NH_2 , alkyl group or phenyl group; and m and n are each a natural number.

[0016] In view of the fact that a phosphorus compound is contained in the composition, the nitrogen content is preferably selected to be 2.5 to 4% by weight per 100 parts by weight of the epoxy resins, curing agent and phosphorus compound combined. If the nitrogen content is less than 2.5% by weight, desired flame retardancy may not be obtained, and if it exceeds 4.0% by weight, the produced composition tends to have too high hygroscopicity.

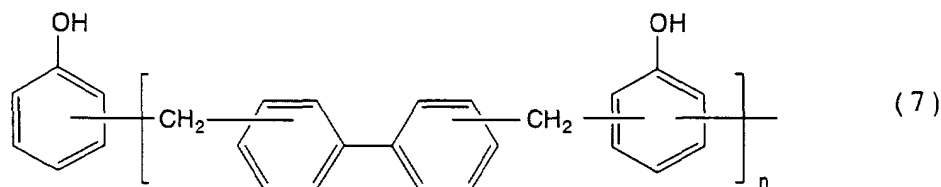
[0017] The phenolic aralkyl resins represented by the following formula (5), the naphthalene aralkyl resins represented by the following formula (6), the biphenyl-modified novolak resins represented by the following formula (7) and the toluene-, xylene- or mesitylene-modified novolak resins represented by the following formula (8) are also preferred for use in the present invention, as these resins have a high content of aromatic rings and are small in ratio of polar groups, and hence high in heat resistance and conducive to the reduction of water absorption of the composition:



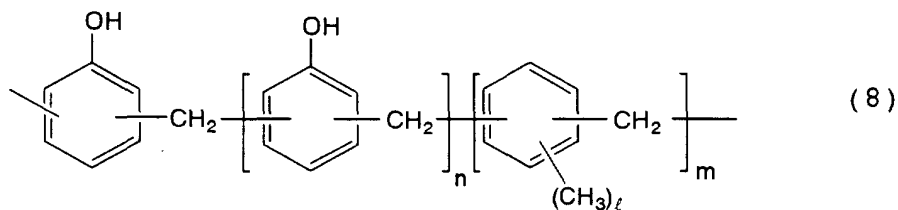
wherein n is a natural number,



wherein n is a natural number,



wherein n is a natural number, and



wherein m and n are each a natural number, and ℓ is a number of 1, 2 or 3.

[0018] In case novolak resins are used as curing agent (B), it is preferable to use a combination of (a) a triazine-modified novolak resin of the formula (4) and (b) at least one member selected from the group consisting of phenolic aralkyl resins of the formula (5), naphthalene aralkyl resins of the formula (6), biphenyl-modified novolak resins of the formula (7) and toluene-, xylene- or mesitylene-modified novolak resins of the formula (8), as this combination of novolak resins has high heat resistance and excellent flame retardancy. The preferred (a)/(b) ratio is 60/40 to 20/80 in view of the above properties.

[0019] When novolak resins are used as a curing agent, the ratio of novolak resins to epoxy resins is preferably 0.8-1.2 in terms of equivalent ratio (phenolic hydroxyl group/epoxy group). If this ratio is outside the above-defined range, the free epoxy groups or phenolic hydroxyl groups remain in the composition to give adverse effects to its heat resistance and water absorption.

[0020] In the present invention, a phosphine oxide compound (C) is used as a phosphorus compound acting as a flame retardant. Phosphine oxide compounds are hardly hydrolyzed and have high chemical resistance. Typical examples of such phosphine oxide compounds are triphenylphosphine oxide and tricresylphosphine oxide. Use of 9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (HCA), a phosphorus compound which, like said phosphine oxide compounds, is hardly hydrolyzed and has high chemical resistance, in combination with said phosphine oxide compound is a preferred embodiment of the present invention.

[0021] In the present invention, the phosphorus content provided by the phosphorus compounds such as phosphine oxide compound (C) is preferably 0.5 - 3% by weight per 100 parts by weight of the epoxy resins, curing agent and phosphorus compounds combined. If the phosphorus content is less than 0.5% by weight, desired flame retardancy may not be obtained, and if it exceeds 3.0% by weight, the produced composition tends to have too high hygroscopicity.

[0022] The flame-retardant resin composition of the present invention comprises as its essential components an epoxy resin, a curing agent and a phosphorus compound functioning as a flame retardant as mentioned above, but it is possible to add other types of epoxy resin and curing agent, or other types of resin and curing agent, curing accelerator, coupling agent or other necessary components within limits not prejudicial to the object of the present invention. In the present invention, however, it is preferred not to add phosphoric esters so as not to deteriorate the innate properties, especially chemical resistance, of the resins and laminates. According to the present invention, excellent flame retardancy can be realized although no phosphoric ester is contained in the composition.

[0023] The flame-retardant resin composition of the present invention finds various ways of utilization. In case the composition is impregnated in a fiber base, it is usually used in the form of a varnish prepared by dissolving the composition in a solvent. As the solvent, there is usually used one having a high capability to dissolve the composition, but use of a poor solvent is not excluded as far as it does not cause any adverse effect.

[0024] The varnish obtained by dissolving the flame-retardant resin composition of the present invention in a solvent may be applied on or impregnated in a base such as woven glass fabric, non-woven glass fabric, or woven or non-woven fabrics made of other materials than glass fiber, and dried at 80 to 200°C to make a prepreg. Such a prepreg may be hot-pressed to produce a laminate or a copper-clad laminate. The flame-retardant resin composition according to the present invention is a thermosetting resin composition having outstanding flame retardancy without containing any halogen compound, and is especially suited for use as a base material of prepregs and laminates for printed circuit boards.

[0025] The present invention will be described in further detail below with reference to the examples thereof.

Example 1

[0026] 100 parts by weight of a phenolic novolak epoxy resin (Epiclon N-770 produced by Dainippon Ink and Chemicals, Inc.), 49 parts by weight of a phenolic aralkyl resin (Milex XLC-LL produced by Mitsui Chemical Co., Ltd.), 31 parts by weight of a triazine-modified phenolic novolak resin (LA-7054 produced by Dainippon Ink and Chemicals, Inc.)

and 40 parts by weight of triphenylphosphine oxide were dissolved in methyl cellosolve to prepare a varnish with a nonvolatile concentration of 60% by weight. The varnish had a phosphorus content of 2.0% by weight and a nitrogen content of 1.7% by weight per 100% by weight of the epoxy resin, curing agent and phosphorus compound combined.

[0027] This varnish was impregnated, in an amount of 80 parts by weight as solids, in 100 parts by weight of a woven glass fabric (0.18 mm thick, manufactured by Nitto Boseki KK) and dried in a 150°C drying oven for 5 minutes to make a prepreg with a resin content of 44.4% by weight.

[0028] 6 sheets of this prepreg were placed one over another, with a 35 µm thick electrolytic copper foil placed on the top and at the bottom of the pile, and subjected to hot compression molding under the conditions of 40 kgf/cm² and 190°C for 120 minutes to obtain a 1.2 mm thick double-side copper-clad laminate.

Examples 2-6 and Comparative Examples 1-4

[0029] The resin compositions of the formulations shown in Table 1 (Examples) and Table 2 (Comparative Examples) were dissolved in methyl cellosolve to prepare varnishes with a nonvolatile concentration of 60% by weight. Thereafter the same procedure as in Example 1 was repeated to make double-side copper-clad laminates.

[0030] Flame retardancy, soldering heat resistance, copper foil peel strength and chemical resistance of the obtained copper-clad laminates were determined. Results of evaluation are shown in Tables 1 and 2. Determination of the above properties was made as described below.

1. Flame retardancy

[0031] This was evaluated by the vertical flame method according to UL-94 standards.

2. Soldering heat resistance

[0032] This was determined according to JIS C-6481. Each test piece, which had been subjected to a moisture absorbing treatment by 2-hour boiling, was immersed in a 260°C solder bath for 120 seconds and examined for abnormality of external appearance.

3. Copper foil peel strength

[0033] This was determined according to JIS C-6481.

4. Chemical resistance

[0034] Per cent loss in weight after 10-minute immersion in a 20 wt% aqueous solution of sodium hydroxide of 80°C (a) and per cent loss in weight after 96-hour immersion in a 20 wt% aqueous solution of potassium persulfate of 70°C (b) were determined.

Table 1

Items		Example 1	Example 2	Example 3	Example 4	Example 5	Example 6
Formulation (part by weight)	Phenolic novolak epoxy resin ¹⁾	100	100	100			100
	Cresol novolak epoxy resin ²⁾						
	Naphthalene aralkyl epoxy resin ³⁾				100		
	Biphenyl-modified epoxy resin ⁴⁾					100	
	Phenolic aralkyl resin ⁵⁾	49	52	39			38
	Triazine-modified phenolic novolak resin ⁶⁾	31	15	23			26
	Phenolic novolak resin ⁷⁾				38		
	Xylene-modified novolak resin ⁸⁾					71	
	HCA ⁹⁾					17	11
	Triphenylphosphine oxide	40	38	35	30	15	20
Properties	Phosphorus (wt%)	2.0	2.1	2.0	2.0	2.0	1.9
	Nitrogen (wt%)	1.7	0.9	1.4	0.0	0.0	1.6
	Flame retardancy (UL94)	V-0	V-0	V-0	V-0	V-0	V-0
	Soldering heat resistance	No abnormality	No abnormality	No abnormality	No abnormality	No abnormality	No abnormality
	Copper foil peel strength (kN/m)	1.5	1.4	1.5	1.3	1.4	1.4
	Chemical resistance (per cent loss in weight)						
	(a) Sodium hydroxide	0.15	0.14	0.13	0.13	0.13	0.15
	(b) Potassium persulfate	0.05	0.05	0.05	0.04	0.03	0.04

Table 2

Formulation (parts by weight)	Items	Comp. Example 1	Comp. Example 2	Comp. Example 3	Comp. Example 4
		100	100	100	100
Properties	Phenolic novolak epoxy resin ¹⁾	32	32	40	38
	Cresol novolak epoxy resin ²⁾	35	37	29	26
	Phenolic aralkyl resin ⁵⁾				
	Triazine-modified phenolic novolak resin ⁶⁾				
Properties	Triphenyl phosphate HCA ⁹⁾		40	40	20 11
	Phosphorus (wt%)	0.0	0.0	1.8	2.0
	Nitrogen (wt%)	2.5	2.6	1.7	1.6
	Flame retardancy (UL94)	HB	HB	V-0	V-0
Properties	Soldering heat resistance	No	No	No	No
	Copper foil peel strength(kN/m)	abnormality	abnormality	abnormality	abnormality
	Chemical resistance (per cent loss in weight)	1.5	1.4	1.4	1.3
	(a) Sodium hydroxide	0.15	0.14	0.98	0.85
Properties	(b) Potassium persulfate	0.02	0.05	0.35	0.25

Notes in Tables 1 and 2:

[0035]

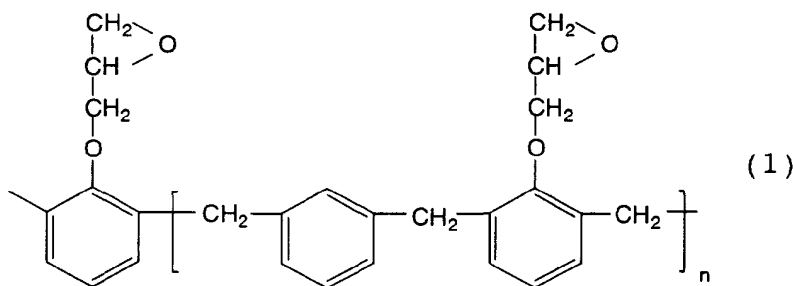
- 1) Epiclon N-770 produced by Dainippon Ink and Chemicals, Inc., epoxy equivalent: 190
- 2) Epiclon N-690 produced by Dainippon Ink and Chemicals, Inc., epoxy equivalent: 210
- 3) ESN-155 produced by Shin Nittetsu Chemical Co., Ltd., epoxy equivalent: 275
- 4) NC-3000P produced by Nippon Kayaku KK, epoxy equivalent: 275
- 5) Milex XLC-LL produced by Mitsui Chemical Co., Ltd., hydroxy equivalent: 175
- 6) LA-7054 produced by Dainippon Ink and Chemicals, Inc., hydroxy equivalent: 125; nitrogen content: 12 wt%
- 7) PR-51470 produced by Sumitomo Durez Co., Ltd., hydroxy equivalent: 105
- 8) R-54537 produced by Sumitomo Durez Co., Ltd., hydroxy equivalent: 190
- 9) 9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide

[0036] As seen from Tables 1 and 2, any of the copper-clad laminates obtained in the Examples has excellent flame retardancy, soldering heat resistance and chemical resistance.

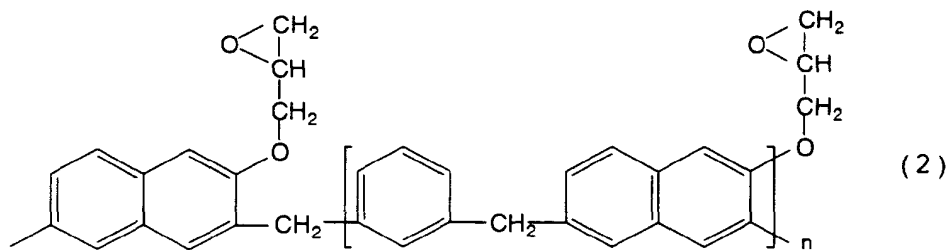
[0037] As described above, the flame-retardant resin composition according to the present invention, without containing any halogen compound, has surpassing flame retardancy and also excels in other properties such as heat resistance. Thus, the present invention provides a novel thermosetting resin composition useful as a halogen-free material which is expected to rise in demand in the future. The prepregs and laminates produced from the above resin composition, and the printed circuit boards comprising such laminates, are also excellent in not only flame retardancy but also other properties such as soldering heat resistance.

Claims

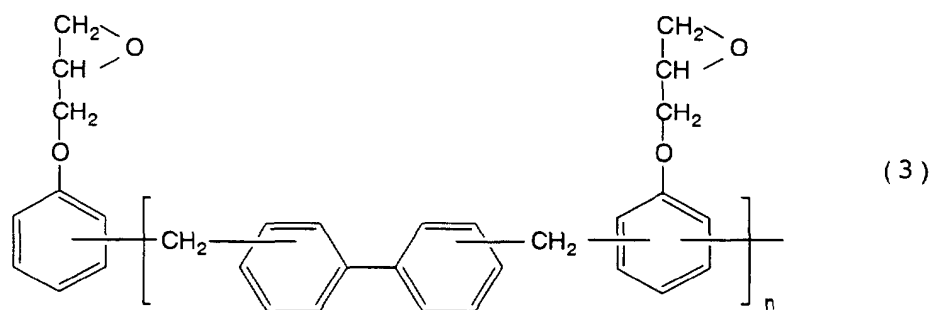
1. A flame-retardant resin composition comprising as essential components (A) an epoxy resin, (B) a curing agent, and (C) a phosphorus compound containing at least as a part thereof a phosphine oxide compound.
2. A flame-retardant resin composition according to Claim 1, wherein part or whole of the epoxy resin (A) is a novolak epoxy resin.
3. A flame-retardant resin composition according to Claim 2, wherein part or whole of the epoxy resin (A) is a phenolic novolak epoxy resin, a cresol novolak epoxy resin, or a mixture of a phenolic novolak epoxy resin and a cresol novolak epoxy resin.
4. A flame-retardant resin composition according to Claim 1, wherein part or whole of the epoxy resin (A) is at least one member selected from the group consisting of phenolic aralkyl epoxy resins represented by the formula (1), naphthalene aralkyl epoxy resins represented by the formula (2) and biphenyl-modified epoxy resins represented by the formula (3):



wherein n is a natural number,

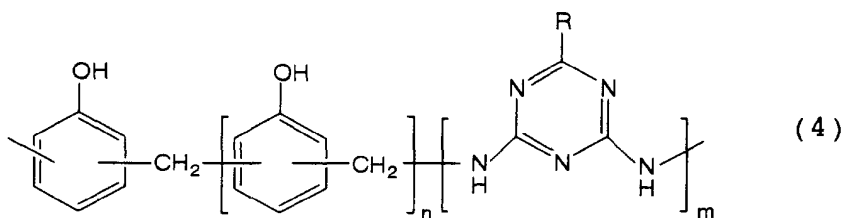


wherein n is a natural number, and



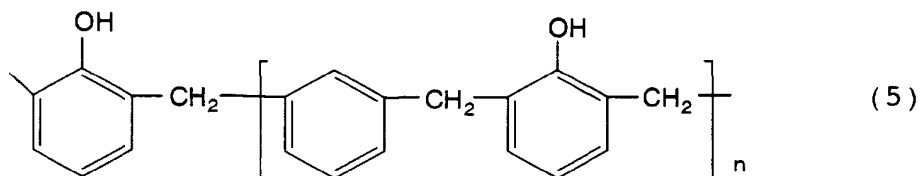
wherein n is a natural number.

- 30
5. A flame-retardant resin composition according to Claim 1, wherein part or whole of the curing agent (B) is a novolak resin.
- 35
6. A flame-retardant resin composition according to Claim 1, wherein part or whole of the curing agent (B) is a triazine-modified novolak resin represented by the formula (4):

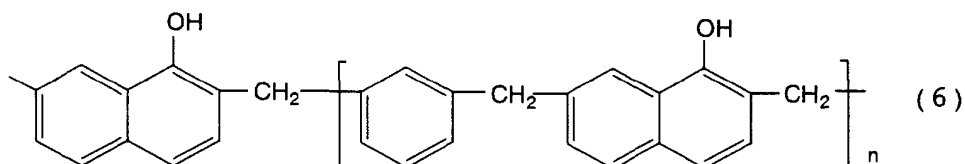


wherein R is NH₂, alkyl group or phenyl group; and m and n are each a natural number.

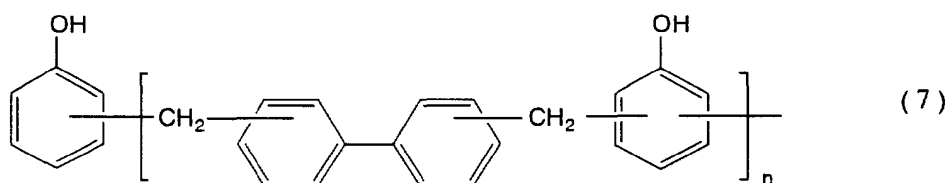
- 50
7. A flame-retardant resin composition according to Claim 1, wherein part or whole of the curing agent (B) is at least one member selected from the group consisting of phenolic aralkyl resins represented by the formula (5), naphthalene aralkyl resins represented by the formula (6), biphenyl-modified novolak resins represented by the formula (7) and toluene-, xylene- or mesitylene-modified novolak resins represented by the formula (8):
- 55



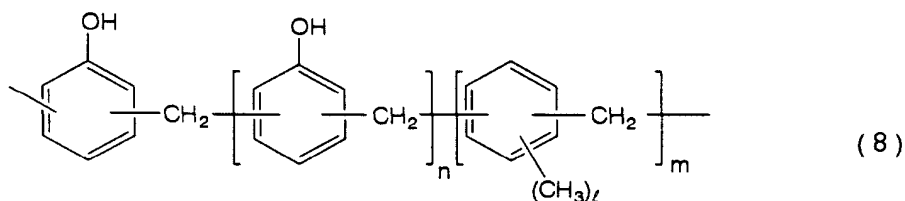
10 wherein n is a natural number,



20 wherein n is a natural number,



30 wherein n is a natural number, and



40 wherein m and n are each a natural number, and ℓ is a number of 1, 2 or 3.

45 **8.** A flame-retardant resin composition according to Claim 1, wherein a triazine-modified novolak resin of the formula (4) and at least one resin selected from phenolic aralkyl resins of the formula (5), naphthalene aralkyl resins of the formula (6), biphenyl-modified novolak resins of the formula (7) and toluene-, xylene- or mesitylene-modified novolak resins of the formula (8) are used in combination as the curing agent.

50 **9.** A flame-retardant resin composition according to any one of Claims 1 to 8, wherein the phosphine oxide compound is triphenylphosphine oxide.

55 **10.** A flame-retardant resin composition according to Claim 9, wherein 9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide is contained as part of the phosphorus compound (C).

11. A prepreg obtained by impregnating a flame-retardant resin composition as set out in any one of Claims 1 to 10 in a fiber base.

12. A laminate characterized in that a single sheet or a pile of two or more sheets of the prepreg as set out in Claim 11 is hot-pressed.

5

13. A copper-clad laminate characterized in that a single sheet or a pile of two or more sheets of the prepreg as set out in Claim 11 is hot-pressed.

10

15

20

25

30

35

40

45

50

55

DERWENT-ACC-NO: 2001-591317

DERWENT-WEEK: 200830

COPYRIGHT 2008 DERWENT INFORMATION LTD

TITLE: Flame retardant resin composition used as base material for prepregs and laminates for printer circuit boards comprises epoxy resin, curing agent and phosphorus compound containing phosphine oxide compound

INVENTOR: TOBASAWA K; TOBISAWA A

PATENT-ASSIGNEE: SUMITOMO BAKELITE CO LTD[SUMB]

PRIORITY-DATA: 2000JP-064370 (March 9, 2000) , 2000JP-007515
(January 17, 2000) , 2000EP-118672 (August 29, 2000)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE
EP 1116774 A2	July 18, 2001	EN
JP 2001200140 A	July 24, 2001	JA
JP 2001254001 A	September 18, 2001	JA
CN 1306041 A	August 1, 2001	ZH
KR 2001076181 A	August 11, 2001	KO
SG 92734 A1	November 19, 2002	EN
US 6551714 B1	April 22, 2003	EN
TW 587094 A	May 11, 2004	ZH
EP 1116774 B1	July 20, 2005	EN
DE 60021344 E	August 25, 2005	DE
CN 1174046 C	November 3, 2004	ZH
DE 60021344 T2	April 27, 2006	DE
JP 4027560 B2	December 26, 2007	JA
KR 701839 B1	March 30, 2007	KO

DESIGNATED-STATES: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT
LU LV MC MK NL PT RO SE SI DE FR GB

APPLICATION-DATA:

PUB-NO	APPL-DESCRIPTOR	APPL-NO	APPL-DATE
EP 1116774A2	N/A	2000EP-118672	August 29, 2000
JP2001200140A	N/A	2000JP-007515	January 17, 2000
JP2001254001A	N/A	2000JP-064370	March 9, 2000
JP 4027560B2	N/A	2000JP-064370	March 9, 2000
TW 587094A	N/A	2000TW-116711	August 18, 2000
US 6551714B1	N/A	2000US-642156	August 21, 2000
SG 92734A1	N/A	2000SG-004757	August 22, 2000
KR2001076181A	N/A	2000KR-050089	August 28, 2000
KR 701839B1	N/A	2000KR-050089	August 28, 2000
DE 60021344E	N/A	2000DE-621344	August 29, 2000
DE 60021344T2	N/A	2000DE-621344	August 29, 2000
EP 1116774B1	N/A	2000EP-118672	August 29, 2000
CN 1306041A	N/A	2000CN-126097	August 31, 2000
CN 1174046C	Previous Publ	2000CN-126097	August 31, 2000

INT-CL-CURRENT:

TYPE	IPC DATE
CIPP	C08J5/24 20060101
CIPP	C08L63/04 20060101
CIPP	C09K21/14 20060101
CIPS	B32B15/08 20060101
CIPS	B32B15/092 20060101
CIPS	B32B15/092 20060101
CIPS	B32B15/20 20060101
CIPS	B32B27/04 20060101
CIPS	C08G59/20 20060101
CIPS	C08G59/24 20060101
CIPS	C08G59/24 20060101
CIPS	C08G59/30 20060101

CIPS	C08G59/32 20060101
CIPS	C08G59/50 20060101
CIPS	C08G59/62 20060101
CIPS	C08G59/62 20060101
CIPS	C08K5/31 20060101
CIPS	C08K5/5397 20060101
CIPS	C08K5/5397 20060101
CIPS	C08K5/5397 20060101
CIPS	C08L63/04 20060101
CIPS	C09K21/12 20060101
CIPS	C09K21/14 20060101
CIPS	H05K1/03 20060101
CIPS	H05K1/03 20060101
CIPS	H05K1/03 20060101
CIPS	H05K3/00 20060101
CIPS	H05K3/00 20060101

ABSTRACTED-PUB-NO: EP 1116774 A2

BASIC-ABSTRACT:

NOVELTY - The flame retardant resin composition comprises an epoxy resin, a curing agent and a phosphorus compound containing at least a phosphine oxide compound.

DESCRIPTION - INDEPENDENT CLAIMS are also included for:

(1) a prepreg obtained by impregnating the resin composition in a fiber base; and

(2) a laminate obtained by hot pressing a single sheet or a pile of two or more sheets of the prepreg.

USE - As base material for prepregs and laminates used for electrical and electronic machine parts, especially printed circuit boards.

ADVANTAGE - The resin composition has excellent flame retardancy. Since

halogen containing flame retardants are not used, toxic halogen compound is not released during pyrolysis. Disposal is made easier. Carbonization of the resin is accelerated and hence combustion is prevented.

EQUIVALENT-ABSTRACTS:

POLYMERS

Preferred Composition: The epoxy resin is a novolak epoxy resin, preferably a phenolic epoxy resin and/or cresol novolak epoxy resin.

The resin is chosen from phenolic aralkyl epoxy resin of formula (I), naphthalene aralkyl epoxy resin of formula (II) and biphenyl-modified epoxy resin of formula (III).

$m, n = \text{natural number.}$

The curing agent comprises phenolic aralkyl resins of formula (V), naphthalene aralkyl resin of formula (VI), biphenyl-modified novolak resin (VII) or xylene- or mesitylene-modified novolak resin of formula (VIII) with novolak resin, preferably triazine-modified novolak resin of formula (IV).

$R = \text{NH}_2, \text{alkyl or phenyl;}$

$l = 1, 2 \text{ or } 3.$

$m, n = \text{natural number.}$

INORGANIC CHEMISTRY

Preferred Composition: The phosphine oxide compound is triphenyl phosphine oxide.

The phosphorus compound comprises also 9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide.

The laminate obtained from the prepreg is a copper-clad laminate.

Phenolic novolak epoxy resin (in parts weight) (100) mixed with phenolic aralkyl resin (49), triazine-modified phenolic novolak resin (31) and triphenyl phosphine oxide (40), was dissolved in methyl cellosolve to obtain a varnish

with 60 weight% non-volatile concentration. The varnish had phosphorus content of 2 wt.% and nitrogen content of 1.7 wt.% with respect to 100 wt.% of the epoxy resin, curing agent and phosphorus compound. The varnish was impregnated on a oven gas fabric and dried at 150 degreesC for 5 minutes to obtain a prepreg with 44.4 wt.% of resin content. 6 prepreg sheets were heat pressed to obtain a laminate with copper cladded on both sides.

The laminate had a flame retardancy of V-0 according to UL-94(TM), no abnormality in soldering heat resistance and copper foil peel strength of 1.5 kN/m. Chemical resistance was excellent with 0.15 wt.% loss when immersed in sodium hydroxide, at 80 degreesC and 0.05 wt.% loss when immersed in potassium persulfate solution, at 70 degreesC.

TITLE-TERMS: FLAME RETARD RESIN COMPOSITION BASE MATERIAL
PREPREG LAMINATE PRINT CIRCUIT BOARD COMPRISE
EPOXY CURE AGENT PHOSPHORUS COMPOUND
CONTAIN PHOSPHINE OXIDE

DERWENT-CLASS: A21 A85 E11 L03 P73 V04 X12

CPI-CODES: A05-A01B; A08-D01; A08-F03; E05-G; E05-G01; E05-G02; L03-H04E1;

EPI-CODES: V04-R07L; V04-X01B;

CHEMICAL-CODES: Chemical Indexing M3 *01* Fragmentation Code B515
B701 B720 B743 B815 B831 G010 G019 G100 M1
M121 M129 M144 M280 M320 M411 M510 M520 M533
M540 M781 Q130 Q140 Q332 Q441 Q454 Q621
Specific Compounds R05423 Registry Numbers 389

Chemical Indexing M3 *02* Fragmentation Code B615
B701 B711 B712 B720 B741 B760 B813 B815 B831
B840 D011 D220 M280 M320 M411 M511 M520 M530
M540 M781 Q130 Q140 Q332 Q441 Q454 Q621 Ring
Index Numbers 40822 Specific Compounds R07571
Registry Numbers 130849

ENHANCED-POLYMER-INDEXING: Polymer Index [1.1] 018 ; G1105 G1092
D01 D19 D18 D31 D50 D76 D86 F31 F30
R00868 181; H0011*R; P0497 P0464
P0226 P0282 M2175 D01 D18;

Polymer Index [1.2] 018 ; G1116*R
G1105 G1092 D01 D11 D10 D19 D18
D31 D50 D76 D87 F31 F30; H0011*R;
P0497 P0464 P0226 P0282 M2175 D01
D18;

Polymer Index [1.3] 018 ; D01 D11 D10
D19 D18 D32 D76 D50 D93 F31 F30;
P0464*R D01 D22 D42 F47; P0453
P0442 P0044 D01 D18 F30; M9999
M2175; M9999 M2200;

Polymer Index [1.4] 018 ; D01 D11 D10
D19 D18 D20 D33 D76 D78 D50 D93 F31
F30; P0464*R D01 D22 D42 F47; P0453
P0442 P0044 D01 D18 F30; M9999
M2175; M9999 M2200;

Polymer Index [1.5] 018 ; ND04; B9999
B4239; K9745*R; K9892; K9789; N9999
N6042*R; N9999 N7192 N7023; Q9999
Q7818*R; K9574 K9483; K9676*R;
K9552 K9483; Q9999 Q7454 Q7330;

Polymer Index [1.6] 018 ; A999 A157*R;
A999 A771;

Polymer Index [1.7] 018 ; D01 D07 D25
D22 D33 D79 D42 D44 D50 D92 F20 P*
5A O* 6A; D01 D19 D18 D33 D50 D76
D93 F20 O* 6A P* 5A R05423 389; A999
A248*R; A999 A771;

Polymer Index [1.8] 018 ; S9999
S1070*R; A999 A419;

Polymer Index [2.1] 018 ; G1105 G1092
D01 D19 D18 D31 D50 D76 D86 F31 F30

R00868 181; G1649*R D01 F07 D11 D10
 D19 D18 D23 D22 D31 D32 D76 D45
 D50 D83 D89 F09 F10; H0033 H0011;
 P0226 P0282*R D01 D18 F30; A999
 A157*R; A999 A782; H0260; P0259*R
 P0226 D01;

Polymer Index [2.2] 018 ; D01 D11 D10
 D19 D18 D32 D76 D50 D93 F31 F30;
 P0453 P0442 P0044 D01 D18 F30; A999
 A157*R; A999 A782;

Polymer Index [2.3] 018 ; D01 D11 D10
 D19 D18 D20 D33 D76 D78 D50 D93 F31
 F30; P0453 P0442 P0044 D01 D18 F30;
 A999 A157*R; A999 A782;

Polymer Index [2.4] 018 ; G1105 G1092
 D01 D19 D18 D31 D50 D76 D86 F31 F30
 R00868 181; G2313 D01 D02 D11 D10
 D19 D18 D31 D76 D50 D87; D01 D02
 D11 D10 D19 D18 D31 D50 D76 D87
 R00862 26; H0033 H0011; P0226
 P0282*R D01 D18 F30; A999 A157*R;
 A999 A782;

Polymer Index [2.5] 018 ; G1105 G1092
 D01 D19 D18 D31 D50 D76 D86 F31 F30
 R00868 181; G2313 D01 D02 G3430
 D11 D10 D19 D18 D31 D50 D76 D88;
 H0033 H0011; P0226 P0282*R D01 D18
 F30; A999 A157*R; A999 A782;

Polymer Index [2.6] 018 ; G1105 G1092
 D01 D19 D18 D31 D50 D76 D86 F31 F30
 R00868 181; G2313 D01 D02 D11 D10
 D19 D18 D31 D76 D50 D89; H0033
 H0011; P0226 P0282*R D01 D18 F30;
 A999 A157*R; A999 A782;

SECONDARY-ACC-NO:

CPI Secondary Accession Numbers: 2001-175470

Non-CPI Secondary Accession Numbers: 2001-440564

